AMENDMENTS TO THE CLAIMS

This listing of claims replaces all previous versions, and listings, of claims pending in this application.

Listing of Claims

1-161. (Canceled)

162. (Currently amended) A compound comprising a peptide chain approximately 12-17 to 40 amino acids in length that binds to G-CSFR and contains a sequence of amino acids selected from the group consisting of GGGLLDICELKLOECARRCN (SEQ ID NO: 209); GRTGGLLDICELKLOECARRCN (SEQ ID NO: 210); LGIEGRTGGGLLDICELKLOECARRCN (SEQ ID NO: 211); LLDICEELKLOEAARRCN (SEQ ID NO: 212); KLLDICELKLOEAARRCN (SEQ ID NO: 213); and LLDICELKLOECARRAN (SEQ ID NO: 343).

of formula (V) (V) $CX^{PL}_{i}X^{PL}_{i}X^{PL}_{i}X^{PL}_{i}X^{PL}_{i}X^{PL}_{i}X^{PL}_{i}X^{PL}_{i}X^{PL}_{i}X^{PL}_{i}C$ (SEQ ID NO: 5) wherein each amino acid is indicated by standard one letter abbreviation, and wherein X^{PL}_{i} is E, G, P, N, R, T, W, S, L, H, A, Q or Y; X^{PL}_{i} is S, T, E, A, D, G, W, P, L, N, V, Y, R or M; X^{PL}_{i} is R, Y, V, Q, E, T, L, P, S, K, M, A or W; X^{PL}_{i} is L, M, G, F, W, R, S, V, P, A, D, C or T; X^{PL}_{i} is V, T, A, R, S, L, W, C, I, E, P, H, F, D or Q; X^{PL}_{i} is E, Y, G, T, Q, M, S, N, A or P; X^{PL}_{i} is C, V, D, G, L, W, E, V, I, S, M or A; X^{PL}_{i} is S, Y, A, W, P, V, L, Q, G, K, F, I, E or D; X^{PL}_{i} is R, W, M, D, H, V, G, A, Q, L, S, E or Y; X^{PL}_{i} is M, L, I, S, V, P, W, F, T, Y, R, or Q; and wherein said compound does not comprise sequence LLDICELKLQECARRCN (SEQ ID NO: 208).

163. (Currently amended) The compound of claim 162, wherein <u>said sequence of amino acids is GGGLLDICELKLQECARRCN (SEQ ID NO: 209).</u>

 $X^{\mu}_{_{4}\text{-}\text{is}} \xrightarrow{E_{_{1}}} X^{\mu}_{_{2}\text{-}\text{is}} \xrightarrow{S\text{ or } \Lambda_{_{1}}} X^{\mu}_{_{3}\text{-}\text{is}} \xrightarrow{R_{_{1}}} X^{\mu}_{_{4}\text{-}\text{is}} \xrightarrow{L_{_{1}}} X^{\mu}_{_{5}\text{-}\text{is}} \xrightarrow{V\text{ or } S_{_{1}}} X^{\mu}_{_{6}\text{-}\text{is}} \xrightarrow{E_{_{1}}} X^{\mu}_{_{5}\text{-}\text{is}} \xrightarrow{S_{_{1}}} X^{\mu}_{_{5}\text{-}\text{is}} X^{\mu}_{_{5}\text{-}\text{is}} \xrightarrow{S_{_{1}}} X^{\mu}_{_{5}\text{-}\text{is}} X^{\mu}_{_{5}\text{-$

164. (Currently amended) The compound of claim 162, wherein <u>said sequence of amino acids is GRTGGLLDICELKLOECARRCN</u> (SEQ ID NO: 210).

the sequence of amino acids is selected from the group consisting of:

GGGLLDICELKLOECARRCN (SEQ ID NO: 209);

GRTGGLLDICELKLOECARRCN (SEQ ID NO: 210);

LGIEGRTGGGLLDICELKLOECARRC- N (SEO ID NO: 211);

LLDICEELKLQEAARRCN (SEQ ID NO: 212); and

KLLDICELKLOEAARRCN (SEO ID NO: 213).

165. (Currently amended) The compound of claim 162, comprising a dimer having the structure of formula (VIII)

$$(\beta A)_{n4} - R^2 - (\beta A)_{n2} \setminus (Lk)_x \setminus (Lk)_y \setminus (\beta A)_{n3} - R^1 - (\beta A)_{n1}$$

wherein R^1 and R^2 are independently selected from the sequences of amino acids of claim 1; formula (V); βA is a β -alanine residue; n1, n2, n3, n4, x and y are independently zero or one with the proviso that the sum of x and y is either one or two; and Lk is a terminal linking moiety selected from the group consisting of a disulfide bond, a carbonyl moiety, a C1-12 linking moiety optionally terminated with one or two-NH-linkages and optionally substituted at one or more available carbon atoms with a lower alkyl substitutent, a lysine residue or a lysine amide.

- 166. (Previously presented) The compound of claim 162, containing a disulfide bond.
- 167. (Previously presented) The compound of claims 162 wherein the N terminus of the peptide is coupled to a polyethylene glycol molecule.
- 168. (Previously presented) The compound of claim 162 wherein the N terminus of the peptide is acetylated.
- 169. (Previously presented) The compound of claim 162, wherein the C terminus of the peptide is amidated.
- 170. (Previously presented) A pharmaceutical composition comprising a therapeutically effective amount of the compound of any claim 162, in combination with a pharmaceutically acceptable carrier.

- 171. (Withdrawn) A method for treating a patient who would benefit from administration of a GCSF modulator, comprising administering to the patient a therapeutically effective amount of the compound of claim 162.
- 172. (Withdrawn) The method of claim 171, wherein the G-CSF modulator is an agonist for the GCSFR.
- 173. (Withdrawn) The method of claim 171, wherein the patient suffers from a depressed neutrophil count.
- 174. (Withdrawn) The method of claim 173, wherein the depressed neutrophil count is associated with a condition selected from the group consisting of chemotherapy-induced neutropenia, AIDS induced neutropenia and community-acquired pneumonia-induced neutropenia.
- 175. (New) The compound of claim 162, wherein said sequence of amino acids is LGIEGRTGGGLLDICELKLOECARRCN (SEQ ID NO: 211).
- 176. (New) The compound of claim 162, wherein said sequence of amino acids is LLDICEELKLQEAARRCN (SEQ ID NO: 212).
- 177. (New) The compound of claim 162, wherein said sequence of amino acids is KLLDICELKLQEAARRCN (SEQ ID NO: 213).
- 178. (New) The compound of claim 162, wherein said sequence of amino acids is LLDICELKLQECARRAN (SEQ ID NO:343).